

Food and Drug Administration  
Rockville MD 20857

Re: GEMZAR™

Docket No. 96E-0314

JUL - 8 1997

Stephen G. Kunin  
Deputy Assistant Commissioner for  
Patent Policy and Projects  
U.S. Patent and Trademark Office  
Box Pat. Ext.  
Assistant Commissioner for Patents  
Washington, D.C. 20231

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JUL 16 1997

PATENT EXTENSION  
A/C PATENTS

Dear Mr. Kunin:

This is in regard to the application for patent term extension for U.S. Patent No. 4,808,614 filed by Eli Lilly & Company under 35 U.S.C. § 156.

A letter sent to you dated March 7, 1997 incorrectly stated in the opening paragraph:

The human drug product claimed by the patent is  
GEMZAR™ (gemcitabine hydrochloride), which was  
assigned New Drug Application (NDA) No. 20-509.

In fact, the statement should have read:

The human drug product identified in the patent extension  
application is GEMZAR™ (gemcitabine hydrochloride),  
which was assigned New Drug Application (NDA) No.  
20-509.

Please let me know if we can be of further assistance.

Sincerely,



Ronald L. Wilson, Director  
Health Assessment Policy Staff  
Office of Health Affairs

cc: Margaret Brumm  
Eli Lilly & Company  
Patent Division/MMB  
Lilly Corporate Center  
Indianapolis, IN 46285